

ABSTRACT OF THE INVENTION

Compounds are disclosed having the general formula R₁-X-R₂, wherein R₁ and R₂ are biologically active groups, at least one of which is polypeptidic. X is a non-peptidic polymeric group. R₁ and R₂ may be the same or different. Preferred R₁ and R₂ groups are interleukin-1 receptor antagonist, 30kDa TNF inhibitor, interleukin-2 receptors and CR1 and muteins thereof. Also included are site selectively modified interleukin-1 receptor antagonist and 30kDa TNF inhibitor.

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